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(54) **Solid polymer electrolyte and use thereof**
Fester Polymerelektrolyt und dessen Verwendung
Electrolyte polymère solide et son utilisation

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Description

[0001] The present invention relates to a crosslinked solid polymer electrolyte which is suitable as a material for electrochemical devices such as battery, capacitor, sensor, condenser and EC (electrochromic) device, and a photo-electric transfer device.

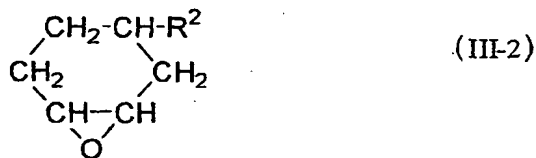
[0002] As an electrolyte constituting an electrochemical device such as a battery, a capacitor and a sensor, those in the form of a solution or a paste have hitherto been used in view of the ionic conductivity. However, the following problems are pointed out. That is, there is a fear of damage of an apparatus arising due to liquid leakage, the mounting and processability of the device are problematic, and subminiaturization and thinning of the device are limited because a separator to be impregnated with an electrolyte solution is required. To the contrary, a solid electrolyte such as inorganic crystalline substance, inorganic glass and organic polymer substance is suggested. The organic polymer substance is generally superior in processability and moldability and the resultant solid electrolyte has good flexibility and bending processability and, furthermore, the design freedom of the device to be applied is high and, therefore, the development thereof is expected. However, the organic polymer substance is inferior in ionic conductivity to other materials at present.

[0003] For example, an attempt that an ethylene oxide-propylene oxide copolymer containing a specific alkali metal salt is utilized as an ion conductive solid electrolyte has already been proposed (JP-A-61-83249, 63-136407 and 2-24975). However, a polymeric substance having more improved ionic conductivity and mechanical properties is still desired.

[0004] It is an object of the present invention to provide a solid polymer electrolyte which has sufficient mechanical strength and flexibility such that conduction is improved and breakage of elements can be prevented, in order to achieve more prevalent application of a solid polymer electrolyte to electrochemical devices.

[0005] This object could be achieved on the basis of the finding that a solid electrolyte having an excellent ionic conductivity and showing little plastic deformation or fluidity even at a high temperature can be obtained by using a multi-component copolymer in which propylene oxide and ethylene oxide are combined with a further crosslinkable oxirane compound and blending this multi-component copolymer with an electrolyte salt compound before or after the crosslinking.

[0006] The present invention is derived from a polyether copolymer having a weight-average molecular weight of 10^5 to 10^7 , which copolymer comprises: (A) 3 to 30 % by mol of repeating units derived from propylene oxide; (B) 96 to 69 % by mol of repeating units derived from ethylene oxide; and (C) 0.01 to 15 % by mol of repeating units derived from a monomer having one epoxy group and at least one reactive functional group represented by formula (III-1) and/or formula (III-2):



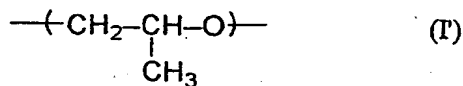
wherein R^1 and R^2 are groups each having a reactive functional group.

[0007] The present invention provides a crosslinked solid polymer electrolyte comprising: (I) a crosslinked material of said polyether copolymer; (II) an electrolyte salt compound; and (III) a plasticizer, which may be added if necessary, selected from an aprotic organic solvent, a derivative or metal salt of linear or branched polyalkylene glycol, and a metal salt of said derivative, where said crosslinked material being produced by utilising reactivity of cross-linkable components of said polyether copolymer.

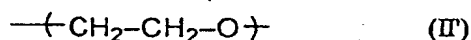
[0008] The present invention also provides a battery comprising said solid polymer electrolyte.

[0009] The copolymer of the present invention has:

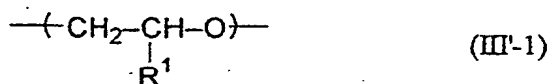
(A) repeating units derived from propylene oxide;



(B) repeating units derived from ethylene oxide; and



(C) repeating units derived from a monomer represented by formula (III-1) or formula (III-2):



and/or



wherein R¹ and R² are groups each having a reactive functional group.

[0010] The crosslinked material of polyether polymer shows excellent shape stability at a high temperature.

[0011] When a plasticizer is mixed into a solid polymer electrolyte, the crystallization of the polymer is suppressed and the glass transition temperature is lowered, achieving a larger magnitude of formation of amorphous phase at a relatively low temperature and thus a higher ion conduction rate. It has been discovered that, when the crosslinked solid polymer electrolyte of the present invention is used, a high-performance battery having a small internal resistance can be obtained. The crosslinked solid polymer electrolyte of the present invention may be a gel. The "gel" means a polymer which has swollen due to absorption of a solvent.

[0012] The polymerization method for obtaining the polyether multi-component copolymer (referred to as the "polyether copolymer" hereinafter) used in the present invention is a polymerization method in which a copolymer is obtained by a ring-opening reaction in the ethylene oxide portion. An example of a polymerization method in which a copolymer is obtained by a ring-opening reaction in the ethylene oxide portion is described in JP-A-62-169823, and JP-A-7-324129.

[0013] Specifically, the polyether copolymer used in the present invention can be obtained by reacting (A) propylene oxide, (B) ethylene oxide and (C) the crosslinkably reactive monomer with each other under the presence or absence of a solvent at a reaction temperature of 10 to 80°C with stirring, using as a ring-opening polymerization catalyst, e.g. a catalyst system mainly containing organic aluminum, a catalyst system mainly containing organic zinc or a catalyst system containing organotin-phosphoric ester condensates.

[0014] Particularly, in case where an oxirane compound having an epoxy group at only both ends is used, when using the organotin-phosphoric ester condensate catalyst, only an epoxy group which does not contain a substituent, i.e. methyl group is used in the polymerization reaction and, therefore, an epoxy group having a methyl group remains in the polymer without being reacted. The organotin-phosphoric ester condensate catalyst is particularly preferable in view of the polymerization degree and properties of the resultant copolymer.

[0015] As the polyether copolymer of the present invention used as a raw material for a crosslinked material, those

comprising 3 to 30 % by mol of the repeating unit (A), 96 to 69 % by mol of the repeating unit (B) and 0.01 to 15 % by mol of the repeating unit (C) are used. Those comprising 5 to 25 % by mol, particularly 10 to 20 % by mol, of the repeating unit (A), 94 to 74 % by mol, particularly 89 to 79 % by mol, of the repeating unit (B) and 0.01 to 10 % by mol, particularly 0.05 to 8 % by mol, of the repeating unit (C) are preferred.

[0016] When the content of the repeating unit (B) exceeds 96 % by mol, crystallization of the oxyethylene chain arises and diffusion transfer of carrier ions is lowered, which results in drastic deterioration of the ionic conductivity of the solid electrolyte. When the content of the repeating unit (B) is smaller than 69 % by mol, dissociation capability of the salt is lowered, which results in deterioration of the ionic conductivity.

[0017] It is generally known that the ionic conductivity is improved by decrease in the crystallizability of polyethylene oxide and decrease in glass transition temperature. It has been found that the effect for improvement of the ionic conductivity is remarkably large by an optimum balance of the monomeric composition of the polyether copolymer of the present invention.

[0018] When the molar ratio of the crosslinking monomer component (monomer capable of forming the repeating unit (C)) is larger than 15 % by mol, the ionic conductivity is drastically lowered and the flexibility of a film is lost, thereby causing problems in processability and moldability.

[0019] The polyether copolymer used in the present invention may be any of a block copolymer and a random copolymer. The random copolymer is preferred because of its large effect of lowering the crystallizability of polyethylene oxide.

[0020] Regarding the molecular weight of the polyether copolymer, the weight-average molecular weight is within a range from 10^5 to 10^7 , and preferably from 5×10^5 to 5×10^6 , so as to obtain excellent processability, moldability, mechanical strength and flexibility. When the weight-average molecular weight is smaller than 10^5 , it is necessary to increase the crosslink density to maintain the mechanical strength and to prevent flow at high temperature and, therefore, the ionic conductivity of the resultant electrolyte is lowered. On the other hand, when it exceeds 10^7 , problems arise in processability and moldability.

[0021] The repeating unit (A) is derived from propylene oxide. The repeating unit (B) is derived from ethylene oxide. The repeating unit (C) is derived from a monomer which has an epoxy group and a reactive functional group represented by the formula (III-1) or the formula (III-2).

[0022] The reactive functional group of the repeating unit (C) is preferably (a) an ethylenically unsaturated group, (b) a reactive silicon group, (c) an epoxy group or (d) a halogen atom.

[0023] The monomer having the ethylenically unsaturated group is preferably an oxirane compound represented by the formula (III-a):



wherein R^3 is a group having an ethylenically unsaturated group.

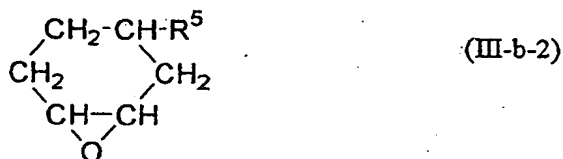
[0024] As the ethylenically unsaturated group-containing oxirane compound, there can e.g. be used allyl glycidyl ether, 4-vinylcyclohexyl glycidyl ether, α -terpinyl glycidyl ether, cyclohexenylmethyl glycidyl ether, p-vinylbenzyl glycidyl ether, allylphenyl glycidyl ether, vinyl glycidyl ether, 3, 4-epoxy-1-butene, 3, 4-epoxy-1-pentene, 4, 5-epoxy-2-pentene, 1, 2-epoxy-5, 9-cyclododecadiene, 3, 4-epoxy-1-vinylcyclohexene, 1, 2-epoxy-5-cyclooctene, glycidyl acrylate, glycidyl methacrylate, glycidyl sorbate, glycidyl cinnamate, glycidyl crotonate and glycidyl-4-hexenoate. Preferable examples thereof include allyl glycidyl ether, glycidyl acrylate and glycidyl methacrylate.

[0025] The monomer having a reactive silicon group, which constitutes the repeating unit (C), is preferably an oxirane compound represented by the formula (III-b-1):



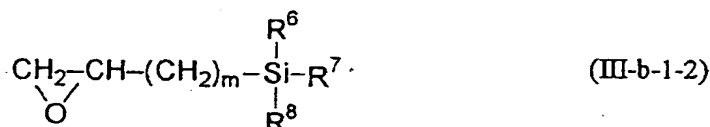
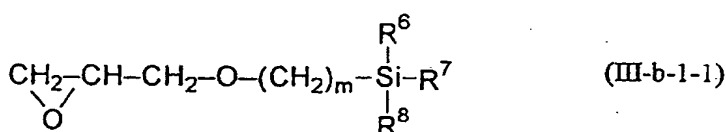
wherein R^4 is a reactive silicon-containing group,

or the formula (III-b-2):

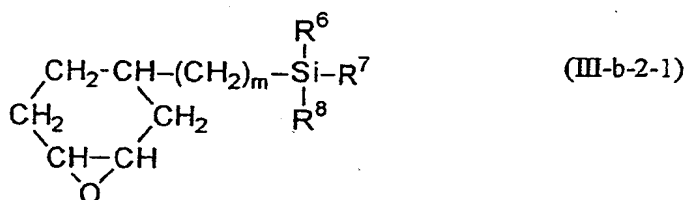


wherein R⁵ is a reactive silicon-containing group.

[0026] The reactive silicon group-containing oxirane compound represented by the formula (III-b-1) is preferably a compound represented by the formula (III-b-1-1) or (III-b-1-2).



[0027] The reactive silicon group-containing monomer represented by the formula (III-b-2) is preferably a compound represented by the formula (III-b-2-1).



[0028] In the formulas (III-b-1-1), (III-b-1-2) and (III-b-2-1), R⁶, R⁷ and R⁸ may be the same or different, but at least one of them represents an alkoxy group and the remainder represent an alkyl group; and m represents 1 to 6.

[0029] Examples of the monomer represented by the formula (III-b-1-1) include 2-glycidyoxyethyltrimethoxysilane, 3-glycidyoxypropylmethyldimethoxysilane, 3-glycidyoxypropyltrimethoxysilane, 4-glycidyoxybutylmethyldimethoxysilane and 4-glycidyoxybutyltrimethoxysilane.

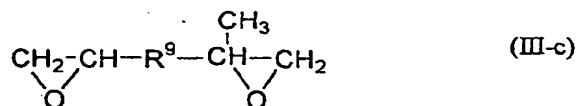
[0030] Examples of the monomer represented by the formula (III-b-1-2) include 3-(1, 2-epoxy) propyltrimethoxysilane, 3-(1, 2-epoxy) propylmethyldimethoxysilane, 3-(1, 2-epoxy) propyldimethylmethoxysilane, 4-(1, 2-epoxy) butyltrimethoxysilane, 4-(1, 2-epoxy) butylmethyldimethoxysilane, 5-(1, 2-epoxy) pentyltrimethoxysilane, 5-(1, 2-epoxy) pentylmethyldimethoxysilane, 6-(1, 2-epoxy) hexyltrimethoxysilane and 6-(1, 2-epoxy) hexylmethyldimethoxysilane.

[0031] Examples of the monomer represented by the formula (III-b-2-1) include 1-(3, 4-epoxycyclohexyl) methyltrimethoxysilane, 1-(3, 4-epoxycyclohexyl) methylmethyl-dimethoxysilane, 2-(3, 4-epoxycyclohexyl) ethyltrimethoxysilane, 2-(3, 4-epoxycyclohexyl) ethylmethyl-dimethoxysilane, 3-(3,4-epoxycyclohexyl) propyltrimethoxysilane, 3-(3,

4-epoxycyclohexyl)propylmethyldimethoxysilane, 4-(3, 4-epoxycyclohexyl) butyltrimethoxysilane and 4-(3, 4-epoxycyclohexyl) butylmethyldimethoxysilane.

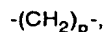
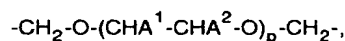
[0032] Among them, 3-glycidoxypropyltrimethoxysilane, 3-glycidoxypropylmethyldimethoxysilane and 4-(1, 2-epoxy) butyltrimethoxysilane are particularly preferable.

[0033] The monomer having the epoxy group as the reactive functional group, which constitutes the repeating unit (C), is preferably an oxirane compound having an epoxy group at both ends represented by the formula (III-c):

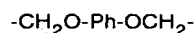


wherein R⁹ is a divalent organic group. R⁹ is preferably an organic group comprising elements selected from hydrogen, carbon and oxygen.

[0034] It is preferable that the group R⁹ in the formula (III-c) is

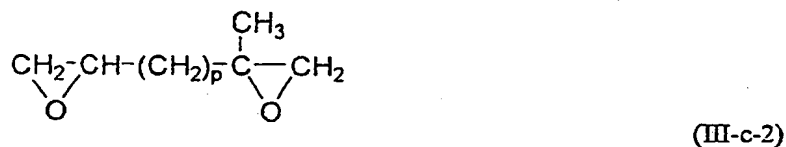
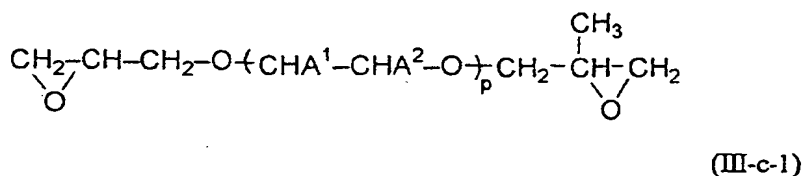


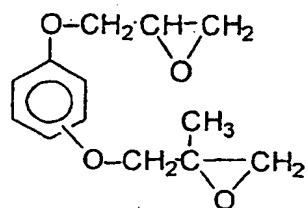
or



wherein A¹ and A² represent hydrogen or a methyl group; Ph represents a phenylene group; and p represents a numeral of 0 to 12.

[0035] The monomer having two epoxy groups at both ends is preferably a compound represented by the following formula (III-c-1), (III-c-2) or (III-c-3):





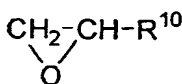
(III-c-3)

[0036] In the above formulas (III-c-1), (III-c-2) and (III-c-3), A¹ and A² represent hydrogen or a methyl group; and p represents a numeral of 0 to 12.

[0037] Examples of the monomer represented by the formula (III-c-1) include 2, 3-epoxypropyl-2', 3'-epoxy-2'-methyl propyl ether, ethylene glycol-2, 3-epoxypropyl-2', 3'-epoxy-2'-methyl propyl ether, and diethylene glycol-2, 3-epoxypropyl-2', 3'-epoxy-2'-methyl propyl ether. Examples of the monomer represented by the formula (III-c-2) include 2-methyl-1, 2, 3, 4-diepoxybutane, 2-methyl-1, 2, 4, 5-diepoxyhexane, and 2-methyl-1, 2, 5, 6-diepoxyhexane. Examples of the monomer represented by the formula (III-c-3) include hydroquinone-2, 3-epoxypropyl-2', 3'-epoxy-2'-methyl propyl ether, and catechol-2, 3-epoxypropyl-2', 3'-epoxy-2'-methyl propyl ether.

[0038] Among them, 2, 3-epoxypropyl-2', 3'-epoxy-2'-methyl propyl ether and ethylene glycol-2, 3-epoxypropyl-2', 3'-epoxy-2'-methyl propyl ether are particularly preferable.

[0039] The monomer having a halogen atom is preferably an oxirane compound represented by formula (III-d):



(III-d)

wherein R¹⁰ is a group having a halogen atom.

[0040] The oxirane compound having a halogen atom includes epichlorohydrin, epibromohydrin and epiodohydrin.

[0041] In the crosslinking method of the copolymer wherein the reactive functional group is an ethylenically unsaturated group, a radical initiator selected from an organic peroxide and an azo compound, or active energy ray such as ultraviolet ray and electron ray can be used. It is also possible to use a crosslinking agent having a silicon hydride.

[0042] As the organic peroxide, there can be used those which are normally used in the crosslinking, such as a ketone peroxide, a peroxy ketal, a hydroperoxide, a dialkyl peroxide, a diacyl peroxide and a peroxy ester. Specific examples of the organic peroxide include 1,1-bis(t-butylperoxy)-3, 3, 5-trimethylcyclohexane, di-t-butyl peroxide, t-butylcumyl peroxide, dicumyl peroxide, 2, 5-dimethyl-2, 5-di(t-butylperoxy)hexane and benzoylperoxide. The amount of the organic peroxide varies depending on the type of the organic peroxide, but it is normally within the range from 0.1 to 10 % by weight based on the whole composition constituting the solid polymer electrolyte and excluding the plasticizer.

[0043] As the azo compound, there can be used those which are normally used in the crosslinking, such as an azonitrile compound, an azoamide compound and an azoamidine compound. Specific examples of the azo compound include 2,2'-azobisisobutyronitrile, 2,2'-azobis(2-methylbutyronitrile), 2, 2'-azobis(4-methoxy-2, 4-dimethylvaleronitrile), 2, 2'-azobis(2-methyl-N-phenylpropionamidine) dihydrochloride, 2, 2'-azobis[2-(2-imidazolin-2-yl) propane], 2, 2'-azobis[2-methyl-N-(2-hydroxyethyl) propionamide], 2, 2'-azobis(2-methylpropane) and 2,2'-azobis [2-(hydroxymethyl) propionitrile]. The amount of the azo compound varies depending on the type of the azo compound, but is normally within the range from 0.1 to 10% by weight based on the whole composition constituting the polymer solid electrolyte and excluding the plasticizer.

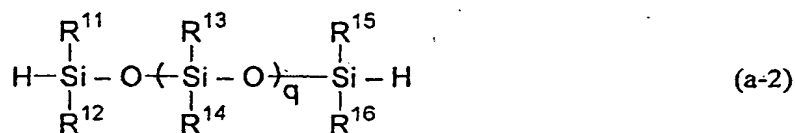
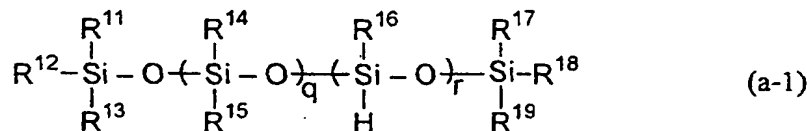
[0044] Particularly preferable examples of the monomer suitable for the crosslinking by active energy ray such as ultraviolet ray include glycidyl acrylate ester, glycidyl methacrylate ester and glycidyl cinnamate ester. Furthermore, as the auxiliary sensitizer, there can be optionally used acetophenones such as diethoxyacetophenone, 2-hydroxy-2-methyl-1-phenylpropan-1-one and phenylketone; benzoin ethers such as benzoin and benzoin methyl ether; benzophenones such as benzophenone and 4-phenylbenzophenone; thioxanthenes such as 2-isopropylthioxanthone and 2, 4-dimethylthioxanthone; and azides such as azidopyrene, 3-sulfonylazidobenzoic acid and 4-sulfonylazidobenzoic acid.

[0045] As a crosslinking aid, there can be optionally used ethylene glycol diacrylate, ethylene glycol dimethacrylate, oligoethylene glycol diacrylate, oligoethylene glycol dimethacrylate, allyl methacrylate, allyl acrylate, diallyl maleate,

triallyl isocyanurate, maleimide, phenylmaleimide and maleic anhydride.

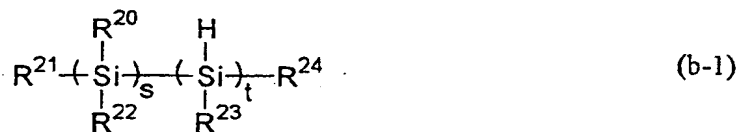
[0046] As a crosslinking agent having a silicon hydride group, which is used for crosslinking the ethylenically unsaturated group, a compound having at least two silicon hydride groups can be used. Particularly, a polysiloxane compound or a polysilane compound is preferable.

[0047] Examples of the polysiloxane compound include a linear polysiloxane compound represented by the formula (a-1) or (a-2), or a cyclic polysiloxane compound represented by the formula (a-3).



[0048] In the formulas (a-1) to (a-3), R^{11} , R^{12} , R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} and R^{19} respectively represent a hydrogen atom or an alkyl or alkoxy group having 1 to 12 carbon atoms; and q and r are an integer provided that $r \geq 2$, $q \geq 0$, $2 \leq q + r \leq 300$. As the alkyl group, a lower alkyl group such as a methyl group and an ethyl group is preferable. As the alkoxy group, a lower alkoxy group such as a methoxy group and an ethoxy group is preferable.

[0049] As the polysilane compound, a linear polysilane compound represented by the formula (b-1) can be used.



[0050] In the formula (b-1), R^{20} , R^{21} , R^{22} , R^{23} and R^{24} respectively represent a hydrogen atom or an alkyl or alkoxy group having 1 to 12 carbon atoms; and s and t are an integer provided that $t \geq 2$, $s \geq 0$, $2 \leq s + t \leq 100$.

[0051] Examples of the catalyst of the hydrosilylation reaction include transition metals such as palladium and platinum or a compound or complex thereof. Furthermore, peroxide, amine and phosphine can also be used. The most popular catalyst includes dichlorobis(acetonitrile)palladium(II), chlorotris(triphenyl-phosphine)rhodium(I) and chloroplatinic acid.

[0052] In the crosslinking method of the copolymer wherein the reactive functional group is a reactive silicon group, the crosslinking can be conducted by the reaction between the reactive silicon group and water. In order to increase the reactivity, there may be used, as a catalyst, organometal compounds, for example, tin compounds such as dibutyltin dilaurate, dibutyltin maleate, dibutyltin diacetate, tin octylate and dibutyltin acetylacetonate; titanium compounds such

as tetrabutyl titanate and tetrapropyl titanate; aluminum compounds such as aluminum trisacetyl acetonate, aluminum trisethyl acetoacetate and diisopropoxyaluminum ethylacetoacetate; or amine compounds such as butylamine, octylamine, laurylamine, dibutylamine, monoethanolamine, diethanolamine, triethanolamine, diethylenetriamine, triethylenetetraamine, cyclohexylamine, benzylamine, diethylaminopropylamine, guanine and diphenylguanine.

[0053] In the crosslinking method of the copolymer wherein the reactive functional group is an epoxy group, e.g. polyamines and acid anhydrides can be used.

[0054] Examples of the polyamines include aliphatic polyamines such as diethylenetriamine, dipropylenetriamine, triethylenetetramine, tetraethylenepentamine, dimethylaminopropylamine, diethylaminopropylamine, dibutylaminopropylamine, hexamethylenediamine, N-aminoethylpiperazine, bis-aminopropylpiperazine, trimethylhexamethylenediamine and dihydrazide isophthalate; and aromatic polyamines such as 4, 4'-diaminodiphenyl ether, diaminodiphenyl sulfone, m-phenylenediamine, 2, 4-toluylenediamine, m-toluylenediamine, o-toluylenediamine and xylylenediamine. The amount of the polyamine varies depending on the type of the polyamine, but is normally within the range from 0.1 to 10% by weight based on the whole composition constituting the solid polymer electrolyte and excluding the plasticizer.

[0055] Examples of the acid anhydrides includes maleic anhydride, dodecenylsuccinic anhydride, chlorendic anhydride, phthalic anhydride, pyromellitic anhydride, hexahydrophthalic anhydride, methylhexahydrophthalic anhydride, tetramethylenemaleic anhydride, tetrahydrophthalic anhydride, methyltetrahydrophthalic anhydride and trimellitic anhydride. The amount of the acid anhydrides varies depending on the type of the acid anhydride, but is normally within the range from 0.1 to 10% by weight based on the whole composition constituting the solid polymer electrolyte and excluding the plasticizer. In the crosslinking, an accelerator can be used. In the crosslinking reaction of polyamines, examples of the accelerator include phenol, cresol, resorcin, pyrogallol, nonyl phenol and 2, 4, 6-tris(dimethylaminomethyl)phenol. In the crosslinking reaction of the acid anhydride, examples of the accelerator include benzyldimethylamine, 2, 4, 6-tris(dimethylaminomethyl)phenol, 2-(dimethylaminoethyl)phenol, dimethylaniline and 2-ethyl-4-methylimidazole. The amount of the accelerator varies depending on the type of the accelerator, but is normally within the range from 0.1 to 10% by weight based on the crosslinking agent.

[0056] In the crosslinking method for crosslinking the copolymer containing a halogen atom, a crosslinking agent such as a polyamine, a mercaptoimidazoline, a mercaptopyrimidine, a thiourea or a polymercaptan can be used. Examples of the polyamine include triethylenetetramine and hexamethylenediamine. Examples of the mercaptoimidazoline include 2-mercaptoimidazoline and 4-methyl-2-mercaptoimidazoline. Examples of the mercaptopyrimidine include 2-mercaptopyrimidine and 4,6-dimethyl-2-mercaptopyrimidine. Examples of the thiourea include ethylenethiourea and dibutylthiourea. Examples of the polymercaptan include 2-dibutylamino-4,6-dimethylcapto-s-triazine and 2-phenylamino-4,6-dimercaptotriazine. The amount of the crosslinking agent to be added may vary depending on the type of the crosslinking agent, but is normally within the range of 0.1 to 30 % by weight based on the whole composition constituting the solid polymer electrolyte and excluding the plasticizer.

[0057] The electrolyte salt compound used in the present invention is preferably soluble in the polyether copolymer or in the crosslinked material of the polyether copolymer. In the present invention, the following electrolyte salt compounds are preferably used.

[0058] That is, examples thereof include a compound composed of a cation selected from metal cation, ammonium ion, amidinium ion and guanidium ion, and an anion selected from chloride ion, bromide ion, iodide ion, perchlorate ion, thiocyanate ion, tetrafluoroborate ion, nitrate ion, AsF_6^- , PF_6^- , stearylsulfonate ion, octylsulfonate ion, dodecylbenzenesulfonate ion, naphthalenesulfonate ion, dodecyl-naphthalenesulfonate ion, 7, 7, 8, 8-tetracyano-p-quinodimethane ion, X^1SO_3^- , $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)\text{N}]^-$, $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)(\text{X}^3\text{SO}_2)\text{C}]^-$ and $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)\text{YC}]^-$, wherein X^1 , X^2 , X^3 and Y respectively represent an electron attractive group. Preferably, X^1 , X^2 and X^3 independently represent a perfluoroalkyl having 1 to 6 carbon atoms or a perfluoroaryl group having 6 to 20 carbon atoms and Y represents a nitro group, a nitroso group, a carbonyl group, a carboxyl group or a cyano group. X^1 , X^2 and X^3 may be the same or different.

[0059] As the metal cation, a cation of a transition metal can be used. Preferably, a cation of a metal selected from Mn, Fe, Co, Ni, Cu, Zn and Ag metals is used. When using a cation of a metal selected from Li, Na, K, Rb, Cs, Mg, Ca and Ba metals, good results are also obtained. Two or more compounds described above may be used as the electrolyte salt compound.

[0060] In the present invention, the amount of the electrolyte salt compound is so that a numeral value of a molar ratio of the number of moles of the electrolyte salt compound to the total number of moles of ether oxygen atom in the main and side chains of the polyether copolymer (the total number of moles of ether oxygen atom included in the polyether copolymer) is preferably within the range from 0.0001 to 5, more preferably from 0.001 to 0.5. When this value exceeds 5, the processability and moldability, the mechanical strength, flexibility and ionic conductivity of the resultant solid electrolyte are deteriorated.

[0061] The plasticizer may be selected from the group consisting of an aprotic organic solvent, a derivative or metal salt of linear or branched polyalkylene glycol, and a metal salt of said derivative.

[0062] The aprotic organic solvent is preferably an aprotic ether or ester. Specific examples thereof include propylene carbonate, γ -butyrolactone, butylene carbonate, ethylene carbonate, dimethyl carbonate, ethylmethyl carbonate, di-

thyl carbonate, 1,2-dimethoxyethane, 1,2-dimethoxypropane, 3-methyl-2-oxazolidone, tetrahydrofuran, 2-methyltetrahydrofuran, 1,3-dioxolane, 4-methyl-1,3-dioxolane, tert-butyl ether, isobutyl ether, 1,2-ethoxymethoxy ethane, ethylene glycol dimethyl ether, ethylene glycol diethyl ether, triethylene glycol dimethyl ether, triethylene glycol diethyl ether, tetraethylene glycol dimethyl ether, tetraethylene glycol diethyl ether, ethylene glyme, ethylene diglyme, methyl tetraglyme, methyl triglyme, methyl diglyme, methyl formate, methyl acetate and methyl propionate. Two or more of the aprotic organic solvent may be used in combination.

[0063] Among the aforementioned examples, propylene carbonate, γ -butyrolactone, butylene carbonate and 3-methyl-2-oxazoline are especially preferable. Triethylene glycol dimethyl ether, triethylene glycol diethyl ether, tetraethylene glycol dimethyl ether and tetraethylene glycol diethyl ether are also especially preferable as the organic solvent.

[0064] The derivative or the metal salt of linear or branched polyalkylene glycol (or a metal salt of said derivative) can be obtained from a polyalkylene glycol having a number-average molecular weight of 200 to 5,000. Examples of the polyalkylene glycol include polyethylene glycol and polypropylene glycol. Examples of the derivatives thereof include an ester derivative and an ether derivative, for example those having an alkyl group having 1 to 8 carbon atoms or an alkenyl group containing 3 to 8 carbon atoms.

[0065] Among the derivatives, examples of the ether derivative include a diether such as dimethyl ether, diethyl ether, dipropyl ether and diallyl ether. Examples of the ester derivative include a diester such as a polyalkylene glycol dimethacrylate ester (e.g. polyethylene glycol dimethacrylate ester), a polyalkylene glycol diacetate ester (e.g. polyethylene glycol diacetate ester), and a polyalkylene glycol diacrylate ester (e.g. polyethylene glycol diacrylate ester).

[0066] Examples of the metal salt include sodium, lithium, dialkyl aluminum salts of polyalkylene glycol.

[0067] Specific examples of the metal salt of the derivative include: sodium, lithium, dialkyl aluminum salts (e.g. dioctyl aluminum salt) of monoethers such as monomethyl ether, monoethyl ether, monopropyl ether, monobutyl ether, monohexyl ether, mono-2-ethyl-hexyl ether, and monoallyl ether; and sodium, lithium, dialkyl aluminum salts of monoesters such as monoacetate ester, monoacrylate ester and monomethacrylate ester. Examples of the metal salt of the polyalkylene glycol derivative include dioctyl aluminum salt of polyethylene glycol monomethyl ether, dioctyl aluminum salt of polyethylene glycol monoethyl ether, and dioctyl aluminum salt of polyethylene glycol monoallyl ether.

[0068] The especially preferable range of the number average molecular weight of the polyalkylene glycol used is from 200 to 2,000.

[0069] The blending ratio of the plasticizer may be changed as desired. The plasticizer in an amount of 0 to 2,000 parts by weight, preferably 1 to 2,000 parts by weight, for example 10 to 1,000 parts by weight, particularly 10 to 500 parts by weight may be added to 100 parts by weight of the polyether copolymer.

[0070] When flame retardancy is required in using the polyether copolymer, the crosslinked material thereof and the solid polymer electrolyte, a flame retardant can be used. That is, an effective amount of those selected from a halide (such as a brominated epoxy compound, tetrabromobisphenol A and chlorinated paraffin), antimony trioxide, antimony pentaoxide, aluminum hydroxide, magnesium hydroxide, phosphate ester, polyphosphate salt and zinc borate as a flame retardant can be added.

[0071] The method for production of the crosslinked solid polymer electrolyte of the present invention is not specifically limited, but the crosslinked solid polymer electrolyte is normally produced by a method of mechanically mixing a polyether copolymer with an electrolyte salt compound or mixing after dissolving them in a solvent, removing the solvent, and crosslinking, or a method of crosslinking a polyether copolymer and mechanically mixing the crosslinked polyether copolymer with an electrolyte salt compound or mixing after dissolving them in a solvent and removing the solvent. As a means for mechanically mixing, various kneaders, open rolls, extruders, etc. can be optionally used. In case of producing the crosslinked solid polymer electrolyte by using the solvent, various polar solvents such as tetrahydrofuran, acetone, acetonitrile, dimethyl formamide, dimethyl sulfoxide, dioxane, methyl ethyl ketone, methyl isobutyl ketone, toluene and ethylene glycol diethyl ether may be used alone or in combination thereof. The concentration of the solution is preferably from 1 to 50 % by weight, but is not limited thereto.

[0072] When the copolymer having an ethylenically unsaturated group is crosslinked by using a radical initiator, the crosslinking reaction is completed at the temperature range of 10 to 200°C within 1 minute to 20 hours. When using energy radiation such as ultraviolet radiation, a sensitizer is normally used. The crosslinking reaction is normally completed at the temperature range of 10 to 150°C within 0.1 second to 1 hour. In case of the crosslinking agent having silicon hydride, the crosslinking reaction can be completed at the temperature of 10 to 180°C within 10 minutes to 10 hours.

[0073] In case that the reactive functional group is a reactive silicon group, the amount of water used in the crosslinking reaction is not specifically limited because the crosslinking reaction easily occurs even in the presence of moisture in the atmosphere. The crosslinking can also be conducted by passing through a cold water or hot water bath for a short time, or exposing to a steam atmosphere.

[0074] When using a polyamine or an acid anhydride in the crosslinking reaction of the copolymer having an epoxy group, the crosslinking reaction is completed at the temperature of 10 to 200°C within 10 minutes to 20 hours.

[0075] The copolymer and crosslinked material of said copolymer shown in the present invention become a precursor

useful as a crosslinked solid polymer electrolyte. The solid polymer electrolyte shown in the present invention is superior in mechanical strength and flexibility, and a large area thin-film shaped solid electrolyte can be easily obtained by utilizing the properties. For example, it is possible to make a battery comprising the solid polymer electrolyte of the present invention. In this case, examples of the positive electrode material include lithium-manganese double oxide, lithium-vanadium double oxide, lithium cobaltate, lithium nickelate, cobalt-substituted lithium nickelate, vanadium pentoxide, polyacene, polypyrene, polyaniline, polyphenylene, polyphenylene sulfide, polyphenylene oxide, polypyrrole, polyfuran, and polyazulene. Examples of the negative electrode material include an interlaminal compound prepared by occlusion of lithium between graphite or carbon layers, a lithium metal and a lithium-lead alloy. By utilizing the high ionic conductivity, the crosslinked solid polymer electrolyte can also be used as a diaphragm of an ion electrode of the cation such as alkaline metal ion, Cu ion, Ca ion and Mg ion.

[0076] The solid polymer electrolyte of the present invention is especially suitable as a material for electrochemical device (e.g. a battery, a capacitor and a sensor).

[0077] The following Examples further illustrate the present invention in detail.

Preparation Example (production of catalyst)

[0078] Tributyltin chloride (10 g) and tributyl phosphate (35 g) were charged in a three-necked flask equipped with a stirrer, a thermometer and a distillation device, and the mixture was heated at 250°C for 20 minutes with stirring under nitrogen stream and the distillate was distilled off to obtain a solid condensate as a residue product. In the following polymerization, this condensate was used as a polymerization catalyst.

[0079] The monomeric composition of the polyether copolymer was obtained by using the element analysis, the iodine value and the ¹H NMR spectrum. A gel permeation chromatography measurement was carried out for measuring the molecular weight of the polyether copolymer. The molecular weight of the polyether copolymer was calculated in terms of the standard polystyrene. The gel permeation chromatography measurement was carried out at 60°C in DMF as the solvent, using the measuring device (RID-6A) manufactured by Shimadzu Corporation and the columns (Showdex KD-807, KD-806, KD-806M and KD-803) manufactured by Showa Denko K.K.

[0080] The glass transition temperature and the heat of fusion were measured, using the differential scanning calorimeter DSC 8230B manufactured by Rigaku Denki K.K., in nitrogen atmosphere and in the temperature range of -100 to 80°C at a temperature rise rate of 10°C/min. The measurement of conductivity σ was carried out by first pinching with platinum electrodes a film which had been vacuum-dried for 72 hours at 30°C under a pressure of at most 133.3 Pa (1 mmHg) and then applying a voltage of 0.5 V and a frequency range of 5 Hz to 13 MHz according to the alternating current method. The conductivity σ was calculated according to a complex impedance method. The flexibility of the solid electrolyte film was evaluated by the presence or absence of breakage when the film having a thickness of 1 mm was bent by 180 degrees at 25°C.

Example 1

[0081] After the atmosphere in a four-necked glass flask (internal volume: 1 L) was replaced by nitrogen, the condensate (300 mg) obtained in the Preparation Example of the catalyst as the catalyst, allyl glycidyl ether (11 g) having a water content adjusted to not more than 10 ppm, propylene oxide (51 g) and n-hexane (500 g) as the solvent were charged in the flask, and ethylene oxide (100 g) was gradually added with monitoring the polymerization degree of propylene oxide by gas chromatography. The polymerization reaction was terminated by using methanol. The polymer was isolated by decantation, dried at 40°C under a normal pressure for 24 hours, and then dried at 45°C under reduced pressure for 10 hours to give 154 g of a polymer.

[0082] The glass transition temperature of this copolymer was -69°C, the weight-average molecular weight was 1,100,000 and the heat of fusion was 5 J/g. The component of propylene oxide was determined by the measurement of NMR spectrum, whereas the component of allyl glycidyl ether was determined by the measurement of the iodine value. The results of the composition analysis (in terms of monomers) are as shown in Table 1.

[0083] The resultant copolymer (1 g) and dicumyl peroxide (0.015 g) as a crosslinking agent were dissolved in acetonitrile (5 ml), and the resultant solution was mixed with lithium perchlorate (electrolyte salt compound) so that a molar ratio of (the number of moles of the electrolyte salt compound to the total number of moles of ether oxygen atoms of copolymer) was 0.05. This mixture solution was cast on a mold made of polytetrafluoroethylene, followed by sufficient drying and further heating under a nitrogen atmosphere at 150°C for 3 hours to give a film. The measurement results of the conductivity and flexibility of the film are shown in Table 1.

Example 2

[0084] Using the monomers shown in Table 1, the copolymerization was conducted by using the same catalyst and

operation as those of Example 1. The resultant polyether copolymer (1 g), triethylene glycol dimethacrylate (0.05 g) and benzoyl oxide (0.015 g) as a crosslinking agent were dissolved in acetonitrile (20 ml), and the resultant solution was mixed with lithium bistrifluoromethanesulfonylimide (electrolyte salt compound) so that a molar ratio of (the number of moles of the electrolyte salt compound to the total number of moles of ether oxygen atoms of copolymer) was 0.05. This mixture solution was heated under a nitrogen atmosphere at 100°C for 3 hours to give a film. The measurement results of the conductivity and flexibility of the film are shown in Table 1.

Example 3

[0085] Using the monomers shown in Table 1, the copolymerization was conducted by using the same catalyst and operation as those of Example 1. The resultant polyether copolymer (1 g), triethylene glycol diacrylate (0.05 g) and 2,2-dimethoxy-1,2-diphenylethan-1-one (0.02 g) as a sensitizing agent were dissolved in acetonitrile (5 ml), and the resultant solution was mixed with lithium perchlorate (electrolyte salt compound) so that a molar ratio of (the number of moles of the electrolyte salt compound to the total number of moles of ether oxygen atoms of copolymer) was 0.05. This mixture solution was cast on a mold made of polytetrafluoroethylene, dried and then exposed to ultraviolet radiation (30 mW/cm², 360 nm) under an argon atmosphere at 50°C for 10 minutes to give a film. The measurement results of the conductivity and flexibility of the film are shown in Table 1.

Example 4

[0086] Using the monomers shown in Table 1, the copolymerization was conducted by using the same catalyst and operation as those of Example 1. The resultant polyether copolymer (1 g) and dibutyltin dilaurate (5 mg) as a catalyst were dissolved in tetrahydrofuran (20 ml) and water (10 μ l) was added, followed by stirring for 15 minutes. After the solvent was removed under normal pressure, the mixture solution was dried at 60°C for 10 hours to give a crosslinked material. The resultant crosslinked material was impregnated with a tetrahydrofuran solution (5 ml) containing lithium perchlorate (100 mg) for 20 hours, heated at 170°C under 80 kgw/cm² for 10 minutes and pressurized to give a film. The measurement results of the conductivity and flexibility of the film are shown in Table 1.

Example 5

[0087] Using the monomers shown in Table 1, the copolymerization was conducted by using the same catalyst and operation as those of Example 1. The resultant polyether copolymer (1 g) and maleic anhydride (150 mg) were dissolved in acetonitrile (10 ml), and the resultant solution was mixed with a solution of lithium perchlorate (electrolyte salt compound) in tetrahydrofuran so that a molar ratio of (the number of moles of the soluble electrolyte salt compound to the total number of moles of ether oxygen atoms of copolymer) was 0.05. This mixture solution was cast on a mold made of polytetrafluoroethylene, dried and then heated at 150°C under 20 kgw/cm² for one hour and pressurized to give a film. The measurement results of the conductivity and flexibility of the film are shown in Table 1.

Example 6

[0088] Using the monomers shown in Table 1, the copolymerization was carried out by using the same catalyst and operation as those of Example 1. The resultant polyether copolymer (1 g) and diethylenetriamine (50 mg) were dissolved in tetrahydrofuran (20 ml), and the reaction was conducted at 40°C for 2 hours. After the solvent was removed under reduced pressure, the mixture solution was dried at 60°C for 6 hours to give a crosslinked material. The resultant crosslinked material was impregnated with a tetrahydrofuran solution (5 ml) containing lithium perchlorate (100 mg) for 20 hours, heated at 160°C under 100 kgw/cm² for 10 minutes and pressurized to give a film. The measurement results of the conductivity and flexibility of the film are shown in Table 1.

Table 1 COPOLYMER AND SOLID POLYMER ELECTROLYTE

	EXAMPLES				
	1	2	3	4	5
Material Monomer charged (mol%)					
Ethylene oxide	70	80	85	90	82
Propylene oxide	27	19	13	9.97	10
Allyl glycidyl ether	3	1			
Glycidyl methacrylate			2		
3-glycidoxypropyltrimethoxy silane				0.03	
2,3-epoxypropyl-2',3'-epoxy-2'-methylpropyl ether					8
Composition of Resulting Copolymer (mol%)					
Ethylene oxide	73	81	87	92.4	83
Propylene oxide	24	18	11	7.57	11
Allyl glycidyl ether	3	1			
Glycidyl methacrylate			2		
3-glycidoxypropyltrimethoxy silane				0.03	
2,3-epoxypropyl-2',3'-epoxy-2'-methylpropyl ether					6
Weight-Average Molecular Weight of Copolymer	1,100,000	2,050,000	2,400,000	3,410,000	2,020,000
Glass Transition Temperature of Copolymer (°C)	-69	-69	-68	-65	-66
Heat of Fusion of Copolymer (J/g)	5	18	43	52	46
Flexibility of Solid Electrolyte Film	No Breakage	No Breakage	No Breakage	No Breakage	No Breakage
Conductivity of Solid Electrolyte Film (S/cm) at 30 °C	6.1 × 10 ⁻³	9.1 × 10 ⁻³	7.9 × 10 ⁻³	6.8 × 10 ⁻³	1.4 × 10 ⁻⁴

Comparative Examples 1-5

[0089] The polyether copolymer shown in Table 2 which was prepared in a manner similar to Example 1 was used.

5 [0090] In Comparative Example 1, the same procedure as that of Example 1 up to the film forming was performed in a manner similar to Example 1 except that only ethylene oxide was polymerized. In Comparative Examples 2 and 5, the film forming was carried out in a manner similar to Example 1. In Comparative Examples 3 and 4, film forming was carried out in a manner similar to Example 2. The results are summarized in Table 2.

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Table 2 COPOLYMER AND SOLID POLYMER ELECTROLYTE

	COMPARATIVE EXAMPLES				
	1	2	3	4	5
Composition of Resulting Copolymer (mol%)					
Ethylene oxide	100	11	74	50	30
Propylene oxide		89	23	48	69.9
Allyl glycidyl ether			3	2	
3-glycidioxypropyltrimethoxy silane					0.1
Weight-Average Molecular Weight of Copolymer	4,200,000	680,000	10,000	860,000	730,000
Glass Transition Temperature of Copolymer (°C)	-57	-65	-67	-67	-65
Heat of Fusion of Copolymer (J/g)	179	0	4	0	0
Flexibility of Solid Electrolyte Film	Breakage Observed	No Breakage	Breakage Observed	No Breakage	No Breakage
Conductivity of Solid Electrolyte Film (S/cm) at 30 °C	5.8×10^{-7}	1.3×10^{-6}	9.2×10^{-6}	9.8×10^{-7}	9.8×10^{-6}

[0091] It is apparent from a comparison of Examples with Comparative Examples that the ionic conductivity and mechanical properties of the crosslinked solid polymer electrolyte formed from the polyether copolymer of the present invention are excellent.

5 Example 7

(1) Production of cathode (positive electrode)

[0092] LiCoO₂ powder (10 g), graphite (KS-15) (7.5 g), the copolymer obtained in Example 3 (7.5 g), dicumyl peroxide (0.025 g), LiBF₄ (0.65 g) and acetonitrile (50 ml) were mixed under stirring by using a disperser to prepare a paste. This paste was coated on an aluminum foil and then dried to adhere a cathode material on the aluminum foil. Then, the cathode material was crosslinked by heating at 150°C for 3 hours in a drier having atmosphere replaced by nitrogen.

(2) Assembling of battery

[0093] A battery was assembled by adhering an Li foil (diameter: 16 mm, thickness: 80 μm) to one main surface of a solid polymer electrolyte film made in Example 2 or 3 and then further the above cathode to the other main surface of the solid polymer electrolyte film. This operation was conducted in a glove box under a dry argon atmosphere.

(3) Charge/Discharge test

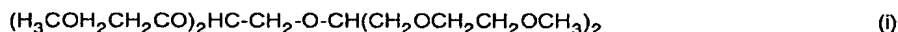
[0094] The resultant battery was charged up to 4.2 V at a temperature of 50°C and a current density of 0.1 mA/cm², and discharged up to 3.0 V. A discharge capacity of 131 mAh per 1 g of LiCoO₂ as an active substance was obtained. The solid polymer electrolyte films of copolymers of Examples 2 and 3 gave the same results.

25 Example 8

1) The cathode was prepared in the same manner as Example 7.

30 2) Assembly of the battery

[0095] 1.6 g of the polyether copolymer obtained in Example 2, 0.4g of a branched polyether (a plasticizer) represented by formula (i):



and 0.02 g of benzoyl peroxide as a crosslinking agent were dissolved in 5 ml of acetonitrile, and then lithium perchlorate was added thereto so that the molar ratio of (the number of moles of soluble electrolyte salt compound)/(the total number of moles of ether oxygen atom in the copolymer) was 0.06. This mixture liquid was cast on a mold made from polytetrafluoroethylene, dried, and then heated in nitrogen atmosphere for 3 hours at 100°C, to obtain a film. The cathode described above, the solid polymer electrolyte film and a Li foil (diameter: 16 mm, thickness: 80 μm) were adhered to assemble a battery. All the operations were carried out in a glove box having a dried argon atmosphere.

45 3) Charging/Discharging Test

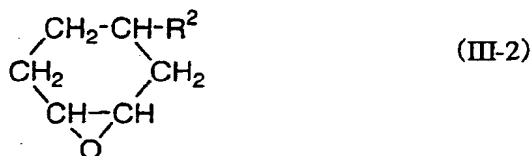
[0096] The resultant battery was charged up to 4.2 V at a temperature of 23°C and a current density of 0.1 mA/cm², and discharged up to 3.0 V. A discharge capacity of 130 mAh per 1 g of LiCoO₂ as an active substance was observed.

[0097] The crosslinked solid polymer electrolyte of the present invention is superior in processability, moldability, mechanical strength, flexibility, heat resistance, etc., and the ionic conductivity is remarkably improved. Accordingly, the crosslinked solid polymer electrolyte of the present invention has an application to electronic apparatuses such as large-capacity capacitor and display device (e.g. electrochromic display) in addition to solid batteries, and an application to antistatic agent for plastic materials.

Claims

1. A solid polymer electrolyte comprising: (I) a crosslinked material of a polyether copolymer; (II) an electrolyte salt compound; and (III) a plasticizer which may be added if necessary and is selected from an aprotic organic solvent, a derivative or a metal salt of a linear or branched polyalkylene glycol, and a metal salt of said derivative, wherein the polyether copolymer has a weight-average molecular weight of 10^5 to 10^7 and comprises:

- (A) 3 to 30 % by mol of repeating unit derived from propylene oxide;
 (B) 96 to 69 % by mol of repeating unit derived from ethylene oxide; and
 (C) 0.01 to 15 % by mol of repeating unit derived from a monomer represented by formula (III-1) and/or formula (III-2):



wherein R^1 and R^2 are groups each having a reactive functional group and the crosslinked material being produced by utilizing reaction of crosslinkable components of the polyether copolymer.

2. The said polymer electrolyte according to claim 1, wherein the reactive functional group in the repeating unit (C) is (a) an ethylenically unsaturated group, (b) a reactive silicon group, (c) an epoxy group or (d) a halogen atom
3. The said polymer electrolyte according to claim 2, wherein the monomer having the ethylenically unsaturated group which constitutes the repeating unit (C) is selected from the group consisting of allyl glycidyl ether, 4-vinylcyclohexyl glycidyl ether, α -terphenyl glycidyl ether, cyclohexenyl methyl glycidyl ether, p-vinylbenzyl glycidyl ether, allyl phenyl glycidyl ether, vinyl glycidyl ether, 3,4-epoxy-1-butene, 3,4-epoxy-1-pentene, 4,5-epoxy-2-pentene, 1,2-epoxy-5,9-cyclododecadiene, 3,4-epoxy-1-vinyl cyclohexene, 1,2-epoxy-5-cyclooctene, glycidyl acrylate, glycidyl methacrylate, glycidyl sorbate, glycidyl cinnamate, glycidyl crotonate and glycidyl-4-hexenoate.
4. The solid polymer electrolyte according to claim 2, wherein the monomer having the reactive silicon group which constitutes the repeating unit (C) is selected from the group consisting of 3-glycidoxy propyl trimethoxy silane, 3-glycidoxy propyl methyl dimethoxy silane, 4-(1,2-epoxy) butyl trimethoxy silane and 2-(3,4-epoxy cyclohexyl) ethyl trimethoxy silane.
5. The said polymer electrolyte according to claim 2, wherein the monomer having two epoxy groups which constitutes the repeating unit (C) is 2,3-epoxypropyl-2',3'-epoxy-2'-methylpropyl ether or ethyleneglycol-2,3-epoxypropyl-2',3'-epoxy-2'-methylpropyl ether.
6. The said polymer electrolyte according to claim 2, wherein the monomer having the halogen atom which constitutes the repeating unit (C) is selected from the group consisting of epichlorohydrin, epibromohydrin and epiiodohydrin.

7. The solid polymer electrolyte according to claim 1, wherein the electrolyte salt compound (II) is a compound composed of a cation selected from metal cation, ammonium ion, amidinium ion and guanidium ion, and an anion selected from chloride ion, bromide ion, iodide ion, perchlorate ion, thiocyanate ion, tetrafluoroborate ion, nitrate ion, AsF_6^- , PF_6^- , stearylsulfonate ion, octylsulfonate ion, dodecylbenzenesulfonate ion, naphthalenesulfonate ion, dodecyl-naphthalenesulfonate ion, 7,7,8,8-tetracyano-p-quinodimethane ion, X^1SO_3^- , $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)\text{N}]^-$, $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)(\text{X}^3\text{SO}_2)\text{C}]^-$ and $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)\text{YC}]^-$ (wherein X^1 , X^2 , X^3 and Y respectively represent an electron attractive group).
8. The solid polymer electrolyte according to claim 7, wherein X^1 , X^2 and X^3 independently represent a perfluoroalkyl group having 1 to 6 carbon atoms or a perfluoroaryl group having 6 to 20 carbon atoms, and Y represents a nitro group, a nitroso group, a carbonyl group, a carboxyl group or a cyano group.
9. The solid polymer electrolyte according to claim 7, wherein the metal cation is a cation of a metal selected from Li, Na, K, Rb, Cs, Mg, Ca, Ba, Mn, Fe, Co, Ni, Cu, Zn and Ag.
10. The solid polymer electrolyte according to claim 1, wherein the aprotic organic solvent is an aprotic organic solvent selected from ethers or esters.
11. The solid polymer electrolyte according to claim 1, wherein the polyalkylene glycol is polyethylene glycol or polypropylene glycol.
12. The solid polymer electrolyte according to claim 1, wherein the derivative of the polyalkylene glycol is an ether derivative or an ester derivative.
13. The solid polymer electrolyte according to claim 1, wherein the metal salt of the polyalkylene glycol is selected from a sodium salt of the polyalkylene glycol, a lithium salt of the polyalkylene glycol, and a dialkyl aluminum salt of the polyalkylene glycol.
14. A battery comprising:
- a solid polymer electrolyte according to claims 1 to 13;
 - a positive electrode; and
 - a negative electrode.

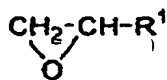
Patentansprüche

1. Fester Polymerelektrolyt umfassend: (I) ein vernetztes Material eines Polyether-Copolymers; (II) eine Elektrolytsalzverbindung; und (III) einen Weichmacher, der bei Bedarf zugegeben werden kann und ausgewählt ist aus einem aprotischen organischen Lösungsmittel, einem Derivat oder einem Metallsalz eines linearen oder verzweigten Polyalkylenglykols, und einem Metallsalz des Derivats, wobei das Polyether-Copolymer ein Gewichtsmittel des Molekulargewichts von 10^5 bis 10^7 aufweist und:

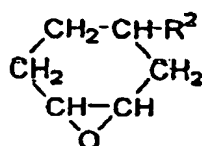
(A) 3 bis 30 Mol-% einer von Propylenoxid abgeleiteten wiederkehrenden Einheit;

(B) 96 bis 69 Mol-% einer von Ethylenoxid abgeleiteten wiederkehrenden Einheit; und

(C) 0,01 bis 15 Mol-% einer von einem Monomer der Formel (III-1) und/oder der Formel (III-2) abgeleiteten wiederkehrenden Einheit:



(III-1)



(III-2)

umfaßt, wobei R¹ und R² Reste sind, die jeweils eine reaktive funktionelle Gruppe aufweisen, und das vernetzte Material unter Ausnutzung der Reaktivität der vernetzbaren Bestandteile des Polyether-Copolymers hergestellt wird.

2. Fester Polymerelektrolyt nach Anspruch 1, wobei die reaktive funktionelle Gruppe in der wiederkehrenden Einheit (C) (a) eine ethylenisch ungesättigte Gruppe, (b) eine reaktive Siliciumgruppe, (c) eine Epoxygruppe oder (d) ein Halogenatom ist.
3. Fester Polymerelektrolyt nach Anspruch 2, wobei das Monomer mit der ethylenisch ungesättigten Gruppe, das die wiederkehrende Einheit (C) darstellt, ausgewählt ist aus Allylglycidylether, 4-Vinylcyclohexylglycidylether, α -Terpenylglycidylether, Cyclohexenylmethylglycidylether, p-Vinylbenzylglycidylether, Allylphenylglycidylether, Vinylglycidylether, 3,4-Epoxy-1-buten, 3,4-Epoxy-1-penten, 4,5-Epoxy-2-penten, 1,2-Epoxy-5,9-cyclododecadien, 3,4-Epoxy-1-vinylcyclohexen, 1,2-Epoxy-5-cycloocten, Glycidylacrylat, Glycidylmethacrylat, Glycidylsorbit, Glycidylcinnamat, Glycidylcrotonat und Glycidyl-4-hexenoat.
4. Fester Polymerelektrolyt nach Anspruch 2, wobei das Monomer mit der reaktiven Siliciumgruppe, das die wiederkehrende Einheit (C) darstellt, ausgewählt ist aus 3-Glycidoxypropyltrimethoxysilan, 3-Glycidoxypropylmethyldimethoxysilan, 4-(1,2-Epoxy)butyltrimethoxysilan und 2-(3,4-Epoxy-cyclohexyl)ethyltrimethoxysilan.
5. Fester Polymerelektrolyt nach Anspruch 2, wobei das Monomer mit zwei Epoxygruppen, das die wiederkehrende Einheit (C) darstellt, 2,3-Epoxypropyl-2',3'-epoxy-2'-methylpropylether oder Ethylenglykol-2,3-epoxypropyl-2',3'-epoxy-2'-methylpropylether ist.
6. Fester Polymerelektrolyt nach Anspruch 2, wobei das Monomer mit dem Halogenatom, das die wiederkehrende Einheit (C) darstellt, ausgewählt ist aus Epichlorhydrin, Epibromhydrin und Epiiodhydrin.
7. Fester Polymerelektrolyt nach Anspruch 1, wobei die Elektrolytsalzverbindung (II) eine Verbindung ist, die aus einem Kation, ausgewählt aus Metallkation, Ammoniumion, Amidinium-Ion und Guanidinium-Ion, und einem Anion, ausgewählt aus Chlorid-Ion, Bromid-Ion, Iodid-Ion, Perchlorat-Ion, Thiocyanat-Ion, Tetrafluorborat-Ion, Nitrat-Ion, AsF₆⁻, PF₆⁻, Stearylsulfonat-Ion, Octylsulfonat-Ion, Dodecylbenzolsulfonat-Ion, Naphthalinsulfonat-Ion, Dodecyl-naphthalinsulfonat-Ion, 7,7,8,8-Tetracyano-pchinodimethan-Ion, X¹SO₃⁻, [(X¹SO₂)(X²SO₂)N]⁻, [(X¹SO₂)(X²SO₂)(X³SO₂)C]⁻ und [(X¹SO₂)(X²SO₂)YC]⁻ (wobei X¹, X², X³ und Y jeweils einen Elektronen anziehenden Rest darstellen) zusammengesetzt ist.
8. Fester Polymerelektrolyt nach Anspruch 7, wobei X¹, X² und X³ unabhängig einen Perfluoralkylrest mit 1 bis 6 Kohlenstoffatomen oder einen Perfluorarylrest mit 6 bis 20 Kohlenstoffatomen darstellen und Y eine Nitrogruppe, eine Nitrosogruppe, eine Carbonylgruppe, eine Carboxylgruppe oder eine Cyanogruppe darstellt.
9. Fester Polymerelektrolyt nach Anspruch 7, wobei das Metallkation ein Kation eines Metalls, ausgewählt aus Li, Na, K, Rb, Cs, Mg, Ca, Ba, Mn, Fe, Co, Ni, Cu, Zn und Ag, ist.
10. Fester Polymerelektrolyt nach Anspruch 1, wobei das aprotische organische Lösungsmittel ein aprotisches organisches Lösungsmittel, ausgewählt aus Ethern oder Estern, ist.
11. Fester Polymerelektrolyt nach Anspruch 1, wobei das Polyalkylenglykol Polyethylenglykol oder Polypropylenglykol ist.
12. Fester Polymerelektrolyt nach Anspruch 1, wobei das Derivat des Polyalkylenglykols ein Etherderivat oder ein Esterderivat ist.

13. Fester Polymerelektrolyt nach Anspruch 1, wobei das Metallsalz des Polyalkylenglykols ausgewählt ist aus einem Natriumsalz des Polyalkylenglykols, einem Lithiumsalz des Polyalkylenglykols und einem Dialkylaluminiumsalz des Polyalkylenglykols.

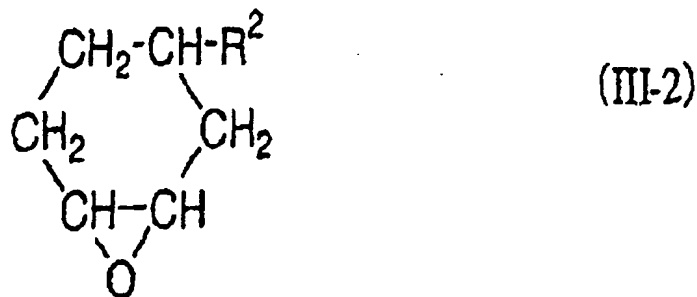
14. Batterie umfassend:

einen festen Polymerelektrolyten nach den Ansprüchen 1 bis 13;
eine positive Elektrode; und
eine negative Elektrode.

Revendications

1. Electrolyte polymère solide comprenant : (I) un matériau réticulé d'un copolymère de polyéther ; (II) un composé salin électrolyte ; et (III) un plastifiant qui peut être ajouté si nécessaire et qui est choisi dans le groupe comprenant un solvant organique aprotique, un dérivé ou un sel métallique d'un polyalkylène glycol linéaire ou ramifié, et un sel métallique dudit dérivé, dans lequel le copolymère de polyéther a un poids moléculaire moyen en poids de 10^5 à 10^7 et comprend :

(A) de 3 à 30 % en mole de motif répétitif dérivé d'oxyde de propylène ;
(B) de 96 à 69 % en mole de motif répétitif dérivé d'oxyde d'éthylène ; et
(C) de 0,01 à 15 % en mole de motif répétitif dérivé d'un monomère représenté par la formule (III-1) et/ou la formule (III-2) :



dans lesquelles R^1 et R^2 sont des groupements ayant chacun un groupement fonctionnel réactif et le matériau réticulé étant produit par utilisation de la réactivité des composants réticulables du copolymère de polyéther.

2. Electrolyte polymère solide selon la revendication 1, dans lequel le groupement fonctionnel réactif du motif répétitif (C) est (a) un groupement éthyléniquement insaturé, (b) un groupement de silicium réactif, (c) un groupement époxy ou (d) un atome d'halogène.
3. Electrolyte polymère solide selon la revendication 2, dans lequel le monomère possédant le groupement éthyléniquement insaturé qui constitue le motif répétitif (C) est choisi dans le groupe comprenant un éther d'allyle et de

glycidyle, un éther de 4-vinyl-cyclohexyle et de glycidyle, un éther de α -terpényle et de glycidyle, un éther de cyclohexényl-méthyle et de glycidyle, un éther de p-vinyl-benzyle et de glycidyle, un éther d'allyl-phényle et de glycidyle, un éther de vinyle et de glycidyle, un 3,4-époxy-1-butène, un 3,4-époxy-1-pentène, un 4,5-époxy-2-pentène, un 1,2-époxy-5,9-cyclododécadiène, un 3,4-époxy-1-vinyl cyclohexène, un 1,2-époxy-5-cyclooctène, un acrylate de glycidyle, un méthacrylate de glycidyle, un sorbate de glycidyle, un cinnamate de glycidyle, un crotonate de glycidyle et un glycidyl-4-hexénoate.

4. Electrolyte polymère solide selon la revendication 2, dans lequel le monomère ayant le groupement silicium réactif qui constitue le motif répétitif (C) est choisi dans le groupe comprenant un 3-glycidoxy propyl triméthoxy silane, un 3-glycidoxy propyl méthyl diméthoxy silane, un 4-(1,2-époxy) butyl triméthoxy silane et un 2-(3,4-époxy cyclohexyl)éthyl triméthoxy silane.
5. Electrolyte polymère solide selon la revendication 2, dans lequel le monomère ayant deux groupes époxy qui constitue le motif répétitif (C) est un 2,3-époxypropyl-2',3'-époxy-2'-méthylpropyl éther ou un éthylèneglycol-2,3-époxypropyl-2',3'-époxy-2'-méthylpropyl éther.
6. Electrolyte polymère solide selon la revendication 2, dans lequel le monomère ayant l'atome d'halogène qui constitue le motif répétitif (C) est choisi dans le groupe comprenant l'épichlorohydrine, l'épibromohydrine et l'épiiodohydrine.
7. Electrolyte polymère solide selon la revendication 1, dans lequel le composé salin électrolyte (II) est un composé formé d'un cation choisi parmi le groupe comprenant un cation métallique, un ion ammonium, un ion amidinium et un ion guanidium, et un anion choisi parmi l'ion chlorure, l'ion bromure, l'ion iodure, l'ion perchlorate, l'ion thiocyanate, l'ion tétrafluoroborate, l'ion nitrate, l' AsF_6^- , le PF_6^- , l'ion sulfonate de stéaryle, l'ion sulfonate d'octyle, l'ion dodécylbenzène sulfonate, l'ion naphthalène sulfonate, l'ion dodécyl-naphthalène sulfonate, l'ion 7,7,8,8-tétracyanop-quinodiméthane, l' X^1SO_3^- , le $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)\text{N}]^-$, le $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)(\text{X}^3\text{SO}_2)\text{C}]^-$ et le $[(\text{X}^1\text{SO}_2)(\text{X}^2\text{SO}_2)\text{YC}]^-$ (dans lesquels X^1 , X^2 , X^3 et Y représentent respectivement un groupe électro-attracteur).
8. Electrolyte polymère solide selon la revendication 7, dans lequel X^1 , X^2 et X^3 représentent indépendamment un groupement perfluoroalkyle ayant de 1 à 6 atomes de carbone ou un groupement perfluoroaryle ayant de 6 à 20 atomes de carbone, et Y représente un groupement nitro, un groupement nitroso, un groupement carbonyle, un groupement carboxyle ou un groupement cyano.
9. Electrolyte polymère solide selon la revendication 7, dans lequel le cation métallique est un cation d'un métal choisi parmi le Li, le Na, le K, le Rb, le Cs, le Mg, le Ca, le Ba, le Mn, le Fe, le Co, le Ni, le Cu, le Zn et l'Ag.
10. Electrolyte polymère solide selon la revendication 1, dans lequel le solvant organique aprotique est un solvant organique aprotique choisi parmi des éthers ou des esters.
11. Electrolyte polymère solide selon la revendication 1, dans lequel le polyalkylène glycol est un polyéthylène glycol ou un polypropylène glycol.
12. Electrolyte polymère solide selon la revendication 1, dans lequel le dérivé du polyalkylène glycol est un dérivé d'éther ou un dérivé d'ester.
13. Electrolyte polymère solide selon la revendication 1, dans lequel le sel métallique du polyalkylène glycol est choisi parmi un sel de sodium du polyalkylène glycol, un sel de lithium du polyalkylène glycol, et un sel d'aluminium dialkyle du polyalkylène glycol.
14. Une batterie comprenant :
 - un électrolyte polymère solide selon les revendications 1 à 13 ;
 - une électrode positive ; et
 - une électrode négative.